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Chapter 1

Nanoscale Thermometry and Temperature Measurement

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1.1 Introduction

Nanoscale temperature measurement concerns the determination of temperature or temperature difference at the sub-micron scale. Applications where it is important to be able to measure local temperature at the nanoscale include microelectronics, optics, microfluidics, chemical reaction and biochemical processes, such as living cells and nano-medicine. Examples of measurement of intracellular temperature fluctuations include the studies by ¹⁻¹³, microcircuit temperature mapping ¹⁴⁻¹⁸, and microfluids ¹⁹⁻²⁴. The topic of temperature measurement at the nanoscale has previously been reviewed ²⁵⁻²⁷. The review ²⁸ provides a general overview of temperature measurement, along with the books ²⁹⁻³⁰. The long standing series ‘Temperature Its Measurement and Control in Science and Industry’ ³¹⁻³⁶ is also worth accessing for its archive on the topic.

In conventional temperature measurement a measurement system typically comprises a transducer to convert a temperature dependent phenomena into a signal that is a function of that temperature, a method to transmit the signal from the transducer, some form of signal processing, a display and method of recording the data. A calibration is then used to convert the measured quantity into a value of temperature and the significance of the temperature can then be considered. Temperature measurement systems can be classified into broad themes, according to the physical contact between the transducer and the medium of interest, Figure 1.1, giving invasive, semi-invasive and non-invasive classifications ^{28, 29}. Examples of each type respectively include thermocouples, thermochromic liquid crystals applied to a surface and observed remotely, and infrared pyrometry.

Figure 1.1: A comparison between the undisturbed medium and a typical temperature measurement application.

Miniaturisation of electronics and optoelectronics with higher switching speeds and associated energy dissipation has increased the importance of a good knowledge of localised heating in order to resolve manufacture and design issues and improve reliability. An accurate knowledge of living cell temperature is proving significant in monitoring cell health, with cancer cells, for example, exhibiting higher temperature than ‘normal tissue’ due to the increased metabolic activity^{37, 5, 38, 7, 8}. Cell temperature alters with cellular activity such as division, gene expression and enzyme reaction, and temperature measurement at this scale can provide valuable diagnostic information.

Temperature can be regarded qualitatively as a measure of hotness or coldness of a body. Quantitatively temperature can be defined from the second law of thermodynamics in terms of the rate of change of entropy with energy²⁹. Thermodynamically temperature is the property that determines whether a system is in thermal equilibrium with another system or systems.

At the nanoscale, issues abound with the definition of temperature, as the scale of the applications means that assumptions of continuum and thermodynamic equilibrium may not hold³⁹. For many practical applications, systems are in a non-equilibrium state and it is difficult to determine the conditions for which equilibrium is established at a local level⁴⁰. A length scale has been proposed³⁹ to identify whether a canonical energy distribution can be assumed. At very low temperatures, e.g. 1 K, a length scale of 0.1 μm was found for Si, providing an indication of the challenge for nanoscale measurements. However the length scale necessary decreases as temperature rises, up to a limit, after which it is reasonably constant. For example a minimum length scale of 100 nm was identified for a temperature of 645 K. The implications of this for nanoscale temperature measurement remain significant, due to non-canonical energy distribution, non-equilibrium and ill-defined material properties. The definition of material properties is particularly important for some transient techniques where knowledge of specific heat capacity is necessary at the temperature of the body concerned.

In order to enable assignment of numerical values to bodies at different temperature, a temperature scale is necessary. The thermodynamic temperature scale is defined by means of perfect heat engines. The SI unit for the thermodynamic temperature scale is the kelvin. As perfect heat engines are not realisable in practice, the International Temperature Scale of 1990, denoted by ITS-90, has been developed as a practical approximation to the thermodynamic temperature scale. The ITS-90 is constructed using a number of overlapping temperature ranges and is believed to represent thermodynamic temperature to within ± 2 mK from 2 K to 273 K, ± 3.5 mK at 730 K and ± 7 mK at 900 K (one standard deviation limits⁴¹). The ITS-90 enables traceability between the measurement that is actually made and the temperature scale defined by the ITS-90.

Due to the delicate and intricate nature of accurate measurements to ITS-90, most practical measurements are made using devices and systems that have been calibrated against

another device which has itself been calibrated using guidelines specified in ITS-90. Calibration in this context is the process of relating the output value from a measurement system to a known input value. This process is illustrated in Figure 1.2 whereby there is a transfer of calibration between practical and specialist devices. The chain between the thermometer or temperature measurement system in use and the ITS-90 may in practice have more links than those shown in Figure 1.2, with each link tending to increase the uncertainty associated with the measurement.

Figure 1.2: A typical chain between practical temperature measurement systems and the International Temperature Scale of 1990.

As well as the physical level of contact between a sensor and the application of interest, thermometric systems can be classified according to whether they are characterised by well-established equations of state that directly relate the measured parameter to temperature, and are known as primary, or if the system requires calibration then it is known as secondary. In addition some materials exhibit an identifiable transition at a specific temperature, an example being phase change, and these can be useful for determining a particular temperature accurately, or to see if a temperature has been exceeded ⁴².

1.2 Methods

A wide range of temperature measurement methods and technologies are available. Conventional technologies such as liquid-in-glass thermometry, thermocouples, and resistance temperature devices (RTDs), have all been adapted to nanoscale applications. Similarly several of the remote observation approaches such as Raman, fluorescence and optical interferometry have been found useful for nanoscale applications. The use of biological agents and cellular features to enable the measurement of temperature in vivo have provoked particular interest. Although a very wide range of temperatures are experienced in nature and science, here the technologies and temperature ranges considered are limited to those addressed by current technologies which are generally applicable to cryogenic temperatures and the temperatures associated with electronic circuit operation.

- Temperature measurement methods relevant to the nanoscale include:
- Liquid and solid-in-tube nanotube thermometry
- Resistance
- Thermocouples, based on the Seebeck effect
- Coulomb blockade
- MEMS (microelectromechanical systems) based resonator quality factor or Fermi-level shift
- Thermochromic liquid crystals, based on crystal phase transitions
- Infrared thermography, based on Planck blackbody emission
- Fluorescence
- Thermo-luminescence or thermographic phosphor thermometry.
- Thermorefectance, based on the temperature dependence of reflection
- Raman, based on inelastic scattering of monochromatic light

- Brownian motion
- Near-field scanning optical microscopy
- Scanning thermal microscopy, using an atomic force microscope with a thermocouple, thermistor or platinum resistance thermometer tip
- Transmission electron microscopy
- Optical interferometry, based on thermal expansion or refractive index change.

In the context of nanoscale applications, the methods can be classified as luminescent and non-luminescent ²⁷. An overview of several of these methods is given in Sections 2.1 to 2.6. The presentation is inevitably brief, but is intended to introduce the principle, provoke interest and offer some relevant references. The Chapters in this book also serve as excellent more in depth treatments.

1.2.1 Infrared

A body emits energy in the form of thermal radiation due to its temperature, with the quantity of radiation rising with increasing temperature. The energy emitted over the electromagnetic spectrum due to temperature by a black body can be modelled by Planck's law. Infrared thermometers measure the thermal radiation emitted by a body due to its temperature and conventional devices have found applications from cryogenic temperatures to over 6000 K. An infrared temperature measurement system generally consists of three elements, the emitting source of interest, the propagation medium, typically space or air, and the measuring device. An infrared measuring device may comprise an optical system, a detector, processing circuit and display ^{29, 43}. The purpose of the optical system is to focus the energy emitted by the target onto the infrared detector, typically converting the energy into an electrical signal. Infrared thermometers can be classified depending on whether the device is

sensitive to all or a specific band of wavelengths, with those that are sensitive to all wavelengths classed as total radiation or broadband thermometers. Devices sensitive to radiation in a specific band of wavelengths are classed spectral band thermometers, ratio thermometers and thermal imagers. Infrared thermography represents a mature technology. For nanoscale applications the spatial resolution is limited, ca. 10 μm , and an estimate of the emissivity is necessary, at the temperature concerned. In addition careful consideration needs to be given to reflection from other sources of thermal radiation in order to ensure that the measurement is close to the temperature of the undisturbed medium, or that appropriate compensation for this is made in the analysis.

1.2.2 Raman Scattering

Raman scattering refers to the inelastic scattering of light. When a molecule is promoted from the ground state to a higher state by incident radiation, it can either return to the original state, which is called Rayleigh scattering, or it can occupy a different vibrational state which is classed as Raman scattering. Raman scattering gives rise to Stokes lines on the observed spectra. Alternatively if a molecule is already in an excited state, it can be promoted to an even higher unstable state and then subsequently return to its ground state. This process is also called Raman scattering and gives rise to anti-Stokes lines on the observed spectra. Monitoring Raman scattering in order to determine temperature involves shining a monochromatic light source, such as a laser, on a sample and detecting the scattered light with a high resolution Raman spectrometer. Fibre optics can be used for remote analysers. The process has the advantage that no preparation of the sample is necessary and small probe volumes, $<1\ \mu\text{m}$, are possible. The method is time consuming and the numbers of samples per second is limited and can be as slow as 0.5 points per second. The application of Raman scattering thermometry has been applied to the polymerase chain reaction ⁴⁴, thermal transport in wires ^{45, 46}, for strain

temperature measurement, temperature and stress ⁴⁷, nanotubes ⁴⁸, nanowires ⁴⁹, and graphene ⁵⁰.

1.2.3 Luminescence

The thermal dependence of phosphor luminescence, band shape, peak energy, intensity, and excited states lifetimes can also be used for temperature measurement. The technique is also known as thermographic phosphor thermometry. A typical set up requires the phosphor or luminescent probe, applied and bonded to a surface, or incorporated within a medium, a light source such as a laser, and a detection system. Types of luminescent thermal probes include organic dyes, ruthenium complexes, quantum dots, Ln^{3+} ions, polymer and organic-inorganic hybrid matrices incorporating emitting centres. The method has the advantages of remote observation, high sensitivity, good spatial resolution and short acquisition periods.

1.2.3.1 Organic Complexes

Organic complexes are readily available and can be selected based on parameters such as the excitation/emission wavelength range required, solubility and facility of vectorisation. An example is nanoparticles of Ru(phen) and PtTFPP embedded in polyacrylonitrile ^{51, 52}. Ru(phen) has a maximum temperature sensitivity of $2.5\% \text{ K}^{-1}$ at 320 K. Alternatively a ratio of intensities from the monomer and excimer/excimerplex can be analysed ⁵³. Similarly the decay of emission lifetimes can be used.

1.2.3.2 Quantum Dots

Quantum dots (QDs) are nanocrystals of a semiconducting material with diameters in the range of about 2 to 10 nm. The high surface-to-volume ratios for these particles results in quantum mechanical properties, such as temperature dependent photoluminescence, which can

be exploited for the purpose of temperature measurement. An example is ZnS coated CdS, which gives a factor of five change in photoluminescence intensity over the range of 100 K to 500 K ^{54, 55}. The method involves observing the QDs through a microscope. A series of factors have limited the use of QDs for thermometry to date, including bluing, bleaching and blinking under continuous illumination ⁵⁶⁻⁵⁸. To overcome some of the practical limitations QDs can be coated with an inert material. The development of an III-nitride quantum dot embedded in a nano-wire for room temperature applications is reported by ⁵⁹. The design of QDs for in-vivo applications has been considered ⁶⁰.

1.2.3.3 Phosphors

The luminescent properties of thermographic phosphors are temperature dependent. This dependency can be observed by applying a thin layer of luminescent material to the system of interest, illumination with UV light and observing the subsequent luminescence. There are two commonly used response modes ^{61, 62}. A pulsed excitation source can be used and after each pulse the phosphor exhibits an exponentially decaying emission, the time constant of which is temperature dependent. Alternatively, pulsed or continuous illumination can be used and the ratio of the emission intensities of two distinct spectral lines can be determined, which is temperature dependent.

Ln^{3+} can be incorporated into a polymer binder and sprayed onto a surface ⁶². A measurement system will typically comprise the surface of interest, the applied coating, a means of exciting the phosphor, such as a laser and optical detector or photomultiplier. The use of a lanthanide coordination polymer $\text{Tb}_{0.957}\text{Eu}_{0.043}\text{cpda}$ as a ratiometric and colorimetric luminescent thermometer for the temperature range 40 to 300 K has been reported ⁶³.

Thin film deposited chromium-doped aluminum oxide ($\text{Cr-Al}_2\text{O}_3$, ruby) thermographic phosphors for temperature measurement have been used ⁶⁴. Zinc silicate ($\text{Zn}_2\text{SiO}_4\text{:Mn}^{2+}$) has

been shown to be highly suitable for fluorescence intensity ratio temperature measurement over the range of 30 to 300 °C with a sensitivity of 12.2% ⁶⁵. The detailed characteristics of fluorescence spectra measurements have been exploited ⁶⁶, in combination with a neural network analysis, in order to determine absolute temperature and have demonstrated this for rhodamine B and copper chloride dyed glycerol. Polydiacetylene (PDA) embedded PVA film can be used to measure thermal gradients effectively ⁶⁷.

1.2.3.4 Biomolecular Intracellular Thermometers

A series of technologies and approaches have been taken to explore possibilities for measuring temperature and temperature at the cellular level. These range from the introduction of fluorescent probes and dyes, the introduction of proteins to control gene expression, the use of materials for the controlled release of chemicals sensitive to temperature, QDs, near infrared imaging, and heating of nano-particles to induce cell response at a known temperature. The use of Temperature-sensitive green emission of fluorescent NaYF₄:Er³⁺,Yb³⁺ nano-particles over the range 25 to 45 °C with application to HeLa cells has been used ⁶⁸. The use of Rhodamine B (RhB) dye employed as a temperature indicator within a single cell has been reported ⁶⁹ with a resolution of better than ± 0.3 °C within the cell cytoplasm, but this reduces to ± 1.8 °C as a result of environmental variations. Temperature sensitive mutants of LacI controlling LacZ expression have been explored ³⁸. Great care is necessary with such approaches as the signal can be a function of duration at a temperature as well as the magnitude of the temperature. Mesoporous materials, such as silica nanoparticles, offer the potential for the controlled release of chemicals that are responsive to temperature changes and this concept has been explored by ⁷⁰⁻⁷². The heating of nanoparticles of gold injected into a cell in combination with optical detection of electron spin resonance in single atomic defects in nitrogen vacancy (NV) diamonds as the sensor has been demonstrated ¹⁰.

An early example ² of using luminescence intensity to measure cell temperature used the temperature dependent luminescence intensity of the $\text{Eu}(\text{tta})_3 \cdot 3\text{H}_2\text{O}$ complex.

A polymer based thermometry solution has been demonstrated ⁵ using PNIPAM combined with the water sensitive fluorophore, DBD-AA. They successfully measured temperatures within the cell over the range 300 to 306 K range with a temperature resolution of 0.3 to 0.5 K and a sensitivity of $S_m = 94\% \text{K}^{-1}$ at 304 K.

Further consideration to the topic of cell based temperature measurement is given in Section 1.2.6.4.

1.2.4 Thermoreflectance

Thermoreflectance exploits the temperature dependence of a material's refractive index, using an optical imaging system to observe the surface of interest, with the temperature profile resolution defined by the spatial resolution of the optical system ^{14, 26}.

1.2.5 Interferometry

Optical interferometry provides both local temperature information, as well as local deformation due to thermal expansion. Interferometers measure differences in optical paths between two light beams, one of which by-passes the test section, while the other passes directly through the test section. A quadriwave lateral shearing interferometer as a wavefront sensor has been used ⁷³ for determining the variation of refractive index for a microwire. Using a water layer on the microwire, and electrical heating, variations in the refractive index generates a distortion of the incident wavefront, enabling mapping of the temperature field.

1.2.6 Non-luminescent

A number of non-luminescent temperature measurement technologies have been developed for nanoscale applications including scanning thermal microscopy, nanolithography, carbon nanotubes, and biomaterials.

1.2.6.1 Scanning Thermal Microscopy

Scanning thermal microscopy uses a small thermocouple with a junction diameter of the order of 20 to 100 nm, formed at a probe tip which is traversed over the surface of interest ⁷⁴⁻⁷⁶. Alternatively an RTD can be mounted on the probe or a combination of a thermocouple and RTD. This can be useful for determining the thermal conductivity of a sample ⁷⁷⁻⁷⁹.

1.2.6.2 Nanolithography

Vapour deposition technology ⁸⁰ can be used to deposit a platinum strip to form a sensor on a surface. Examples include decomposition of organic inks such as $(\text{CH}_3)_3\text{Pt}$. Combinations of materials can be built up by appropriate layering in order to ensure electrical isolation and lead-outs. US Patent 6905736 ⁸¹ reports a Pt(Ga)-W(Ga) junction with a temperature coefficient of 5.4 mV/°C. 100 nm² gold-nickel thermocouples on silicon and 500 nm² thermocouples on quartz have been produced ⁸². A challenge with vapour deposition sensors is ensuring robustness of the sensor, and chemical consistency. It is well known that some materials aggressively leach elements from a substrate, gradually degrading the performance of a sensor with time.

1.2.6.3 Carbon Nanotube Thermometry

Liquid in glass thermometry provided the basis for modern temperature measurement ²⁹, ³⁰. Carbon nanotube thermometry was originally proposed ⁸³ using gallium as the thermometric

liquid, with a scanning electron microscope to observe the meniscus. The coefficient of thermal expansion of gallium has been observed⁸³ to vary linearly over the range 323 K to 773 K, with values close to the macroscopic quantities. Examples of other thermometric liquids proposed and demonstrated for difference temperature ranges include Pb using a ZnO nanotube⁸⁴, Au(Si) using a gallium oxide nanotube⁸⁵, and In using In₂O₃ nanotubes⁸⁶. The use of carbon nanotubes to provide good thermal contact between a microscope and sample has been reported⁸⁷.

1.2.6.4 Biomolecular Based Thermometry

Many molecules and biological components exhibit temperature dependent behaviours that can be exploited for temperature measurement on the nanoscale. Examples include the introduction of proteins to control gene expression, crystal defects, luminescence.

As noted previously, temperature represents a fundamental parameter for the operation of living cells and a number of mechanisms are apparent that can be exploited or mimicked in the measurement or control of temperature⁸⁸. For histidine kinase, DesK, from *Bacillus subtilis*,⁸⁹ report how the plasticity of the helical domain influences the catalytic activities of the protein, through conformational signals transmitted by membrane connections enabling interhelical rearrangements, controlling the alternation between output autokinase and phosphatase activities. Of interest⁸⁹ suggest that the structural comparison of the different DesK variants indicates that incoming signals can take the form of helix rotations and asymmetric helical bends comparable to some other sensing systems and that similar switching mechanism could be operational in a wide range of sensor histidine kinases, or indeed exploited in new designed mechanisms.

Crystal defects, notably the nitrogen vacancy (NV) centre, in diamond can be exploited to enable the measurement of a series of physical parameters including temperature. The optical

excitation of NV nanodiamonds with 532 nm laser light and fluorescence collection with a confocal microscope is reported by ¹⁰, yielding a noise floor of 130 mK/Hz^{0.5} and accuracies as good as 1 mK. Embedded in nanocrystals they can provide robust single-photon sources and fluorescent biomarkers ¹³. Nanodiamonds can be injected into a cell, and excited with a laser to alter their spinstate and the light emitted measured. By using more than one nanodiamond the temperature difference within a single cell can be evaluated as reported by ¹¹, detecting temperature differences as small as 1.8±0.3 mK. Interestingly ¹¹ report that by combining nanodiamonds with laser heating of gold particles also injected into a cell, the temperature of the cell can be controlled.

Living cells sense temperature through proteins, nucleic acids and mRNAs that either modify their conformation structure directly ⁹⁰⁻⁹² or undergo complex reactions. Cells respond to harmful temperature changes by inducing heat shock or cold shock proteins. RNA temperature sensing capability is located in specific areas of RNA structure, especially the 5' untranslated region of certain heat shock and virulence genes, which shields the ribosome-binding sites at physiological temperatures ⁹³. Under the onset of heat shock, destabilisation of RNA commences with associated release of ribosome-binding sites and translation initiation. For cold-shock, cell growth tends to stop, accompanied by a dramatic reduction in bulk gene expression and cold shock proteins are expressed. The underlying mechanism in cells involves a gradual melting of a weak stem-loop structure rather than a switch between two mutually exclusive conformations. As such RNA thermometers can be considered as 'molecular dimmers' rather than discrete on and off state switches ⁹³. Naturally occurring thermometers represent highly complex systems, but simpler RNA stem loop structures, see for example Figure 1.3, containing the ribosome binding site can be produced ^{94, 95, 93}. Ideas for the use of the DNA backbone for temperature sensors and switches have been explored ¹².

Figure 1.3: Synthetic RNA thermometer example. After ⁹⁴.

1.3 Selection

Conventionally, the selection of a specific sensor or system for temperature measurement involves consideration of a number of parameters including:

- uncertainty;
- temperature range;
- thermal disturbance;
- level of contact;
- size of the sensor;

- transient response;
- sensor protection;
- availability;
- cost.

In the case of nanoscale applications, however, the small scale and specific requirements, along with available technology may dictate the use of a particular method. Despite any apparent lack of choice, consideration still needs to be given to the uncertainty associated with the measurement. A common assumption is that because fundamental physics is being used, there is no need to consider a measurement uncertainty. However reasonably simple analysis approaches can be used to evaluate the thermal disturbance associated with the incorporation of a probe into a medium of interest ²⁹. This may, depending on the application, involve just a conduction equation analysis accounting for any differences in thermal conductivity between the undisturbed medium, and that with the sensor embedded, or coating applied, or may involve a thermal radiation calculation to evaluate whether the presence of a probe has altered the thermal radiation exchanges significantly. It is all too easy to alter the temperature of a sample inadvertently due to the presence of an optical head. In addition new methods such as the observation of Brownian motion or nanoparticles are emerging ⁹⁶, providing new opportunities and technology options to assess.

A summary of the approximate capabilities of a range of methods is given in Table 1.1. This has been adapted from the information given in ^{27, 97, 26, 28}.

Method	Spatial Resolution (μm)	Temperature resolution (K)	Response time (μs)
Infrared thermography	10	10^{-1}	10
Thermoreflectance	10^{-1}	10^{-2}	10^{-1}
Raman	1	10^{-1}	10^6
Thermocouple	10^2	10^{-1}	10
Fluorescence	10^{-1}	10^{-2}	10
Near-field scanning optical microscopy	10^{-2}	10^{-1}	10
Liquid crystals	10	10^{-1}	10^2
Scanning thermal microscopy	10^{-1}	10^{-1}	10^2
Transmission electron microscopy	10^{-2}	10^{-1}	10
Optical interferometry	1	10^{-5}	10^{-3}

Table 1.1: Typical spatial resolution, temperature resolution and response times for a range of nanoscale relevant temperature measurement methods.

In assessing the performance of a luminescent thermometer a useful parameter is the relative sensitivity, defined by

$$S = \frac{\left(\frac{\partial Q}{\partial T}\right)}{Q} \quad (1.1)$$

Where Q , is an experimental parameter and corresponds to the quenching of luminescence with temperature. For example ⁹⁸ determined a sensitivity of 4.9% K^{-1} for orthosilicate/aminopropyltriethoxysilane (TEOS/APTES) organosilica shell and co-doped with $[\text{Eu}(\text{btfa})_3(\text{MeOH})(\text{bpeta})]$ and $[\text{Tb}(\text{btfa})_3(\text{MeOH})(\text{bpeta})]$ β -diketonate chelates using the integrated areas I_{Eu} and I_{Tb} of the $^5\text{D}_0\text{--}^7\text{F}_2$ (I_{Eu} at 612 nm) and $^5\text{D}_4\text{--}^7\text{F}_5$ (I_{Tb} at 545 nm) transitions, in this case, $Q = I_{\text{Eu}}^2 - I_{\text{Tb}}^2$. Spatial and temporal resolutions of 0.42 μm and 4.8 ms respectively have been reported ⁹⁹.

Values for sensitivity for a range of luminescent molecular thermometers are given in Table 1.2, adapted from ²⁷.

Phosphor	S_m	ΔT (T_m)
Ru-phen	2.5	280-325 (320)
Fluorescein D-Texas-Red A in DNA	4.5	278-318 (295)
PtOEP	4.6	290-320 (305)
RhB in PDMS	2.3	293-373 (363)
C70 in PtBMA	2.2	293-363 (363)
PNIPAM	10.4	293-318 (308)
Mutat of lacI (Its265)	19.6	308-318 (318)
(CdSe)ZnS QDs	2.2	278-313 (313)
YAG:Ce nano-particles	0.2	315-350 (350)
$Y_2O_3:Eu^{3+}$	2.6	473-973 (973)
Eu(tta) ₃ .3H ₂ O embedded into PMMA	4.4	293-333 (330)

Table 1.2: Figure of merit sensitivity and temperature range for a selection of luminescent molecular thermometers. Data adapted from ²⁷.

1.4 Conclusions

The measurement of temperature at the nanoscale represents a practical reality with a wide range of techniques having been demonstrated. A particular challenge remains for systems where thermal gradients exist, the temperature is at cryogenic levels and the thermal mass is low, arising from the fundamental requirement for sufficient continuum to enable a meaningful definition of temperature for the system concerned. Analysis methods have been proposed to enable calculation of the necessary length scales for meaningful measurements of temperature, with the length scale dependent on material properties and the temperature. The host of technologies available have now led to choice for some applications as well as bespoke solutions emerging, for example for the measurement of temperature on micro-elements in circuitry and within cells. The development of understanding at the cellular level offers opportunities for the transfer of this knowledge to the design of new synthetic devices for both measurement and control. The pioneering work on nanoscale temperature measurement has

certainly opened up insight across many domains, as well as posing some fundamental challenges both in terms of the technology and making sense of the data and its validity.

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